### **PATENT COOPERATION TREATY**

From the INTERNATIONAL SEARCHING AUTHORITY

То:			PCT			
see form PCT/ISA/220			WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (PCT Rule 43bis.1)			
			(day/month/year) see form PCT/ISA/210 (second sheet)			
Applicant's or agent's file reference see form PCT/ISA/220			FOR FURTHER A See paragraph 2 belo	10.10.1		
International application No. PCT/EP2004/011386		International filing date (c 07.10.2004	Priority date (day/month/year) 09.10.2003			
International Patent Classifica C07D277/56, A61K31/4			and IPC			
Applicant SMITHKLINE BEECHAM CORPORATION						
1. This opinion contains indications relating to the following items:  □ Box No. I □ Basis of the opinion □ Box No. II □ Priority □ Box No. III □ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability □ Box No. IV □ Lack of unity of invention □ Box No. V □ Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement □ Box No. VI □ Certain documents cited □ Box No. VII □ Certain defects in the international application □ Box No. VIII □ Certain observations on the international application						
2. FURTHER ACTION If a demand for international preliminary examination is made, this opinion will usually be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA"). However, this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notifed the International Bureau under Rule 56.1bis(b) that written opinions of this International Searching Authority will not be so considered. If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of three months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later. For further options, see Form PCT/ISA/220.						
3. For further details, see notes to Form PCT/ISA/220.						

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## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/EP2004/011386

1.		egard to the language, this opinion has been established on the basis of the international application in iguage in which it was filed, unless otherwise indicated under this item.				
	la	nis opinion has been established on the basis of a translation from the original language into the following nguage , which is the language of a translation furnished for the purposes of international search nder Rules 12.3 and 23.1(b)).				
2.		egard to any nucleotide and/or amino acid sequence disclosed in the international application and sary to the claimed invention, this opinion has been established on the basis of:				
	a. type	e of material:				
		a sequence listing				
		table(s) related to the sequence listing				
	b. format of material:					
		in written format				
		in computer readable form				
	c. time	time of filing/furnishing:				
		contained in the international application as filed.				
		filed together with the international application in computer readable form.				
		furnished subsequently to this Authority for the purposes of search.				
3.	h C	n addition, in the case that more than one version or copy of a sequence listing and/or table relating ther as been filed or furnished, the required statements that the information in the subsequent or additional opies is identical to that in the application as filed or does not go beyond the application as filed, as ppropriate, were furnished.				

## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/EP2004/011386

	Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability					
			ntion appears to be novel, to involve an inventive step (to be non nave not been examined in respect of:			
	the entire international application,					
$\boxtimes$	claims Nos. 7,8					
beca	ause:					
☒	the said international application, or the said claims Nos. 7,8 relate to the following subject matter which does not require an international preliminary examination (specify):					
	see separate sheet					
	the description, claims or drawings (indicate particular elements below) or said claims Nos. are so unclear that no meaningful opinion could be formed (specify):					
	the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.					
	no international search report has been established for the whole application or for said claims Nos.					
	the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:					
	the written form		has not been furnished			
			does not comply with the standard			
	the computer readable form		has not been furnished			
			does not comply with the standard			
	the tables related to the nucleotide and/or amino acid sequence listing, if in computer readable form only, or not comply with the technical requirements provided for in Annex C-bis of the Administrative Instructions.					
	See separate sheet for further of	detai	ls .			

## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/EP2004/011386

Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

1-8

No: Claims

Inventive step (IS)

Yes: Claims

No: Claims

1-8

Industrial applicability (IA)

Yes: Claims

1-6

No: Claims

2. Citations and explanations

see separate sheet

#### III. Non-establishment of opinion

Claims 7 and 8 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(i) PCT).

#### V. Reasoned statement

Reference is made to the following documents:

D1: WO02/096895

#### **Novelty**

Although general formula (I) of D1 generically encompasses the present compounds, there is no specific disclosure of compounds wherein the substituent of the R<sup>6</sup> phenyl group is at the 4-position and is isopropyl. The present compounds may therefore be considered a novel selection from D1. Claims1-8 fulfil the requirements of Article 33(2) PCT.

#### Inventive step

According to the present application, the technical problem to be solved is the provision of a selective dual hPPAR alpha/gamma agonist. Such an agonist is defined as having an EC50 for PPAR-alpha and PPAR-gamma which is at least 10 fold lower than that for PPAR-delta.

Both of the tested compounds of D1 fulfil the desired criteria, as shown by the table on p. 25 of this document. It would be obvious to solve the above-formulated technical problem by providing a further compound falling within the generic definition of D1, formula (I), and in particular to provide a compound with very close structural similarity to the tested examples of D1. Solving the problem by replacing the 4-ethyl substituent of D1, example 2 by a 4-isopropyl substituent cannot be considered inventive.

Claims 1-8 do not fulfil the requirements of Article 33(3) PCT.

#### Industrial applicability

Claims 1-6 fulfil the requirements of Article 33(4) PCT.

No unified criteria exist in the PCT Contracting States for assessing whether present claims 7 and 8 are industrially applicable. The patentability can be dependent upon the formulation of the claims. For example, the EPO does not

# WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (SEPARATE SHEET)

International application No.

PCT/EP2004/011386

consider claims to the use of a compound in medical treatment to be industrially applicable, but allows claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.